

0/777,244

(FILE 'HOME' ENTERED AT 21:50:37 ON 10 DEC 2004)

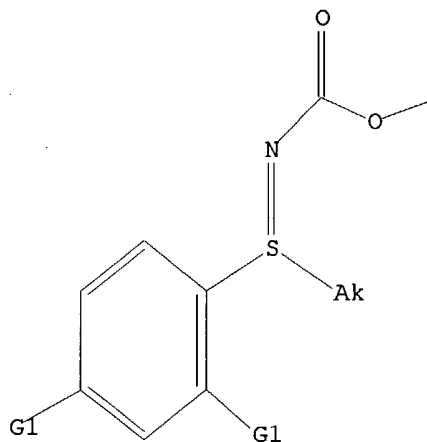
FILE 'REGISTRY' ENTERED AT 21:51:24 ON 10 DEC 2004

L1 STRUCTURE UPLOADED

=> D L1

L1 HAS NO ANSWERS

L1 STR



G1 MeO, EtO, n-PrO, i-PrO, n-BuO, i-BuO, s-BuO, t-BuO, NO2, H, X

Structure attributes must be viewed using STN Express query preparation.

=> S L1

SAMPLE SEARCH INITIATED 21:51:49 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 369 TO ITERATE

100.0% PROCESSED 369 ITERATIONS

3 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*

BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 6228 TO 8532

PROJECTED ANSWERS: 3 TO 162

L2 3 SEA SSS SAM L1

=> S L1 FULL

FULL SEARCH INITIATED 21:51:53 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 7654 TO ITERATE

100.0% PROCESSED 7654 ITERATIONS

56 ANSWERS

SEARCH TIME: 00.00.01

L3 56 SEA SSS FUL L1

=> FIL CAPLUS

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

155.42

155.63

FILE 'CAPLUS' ENTERED AT 21:51:58 ON 10 DEC 2004

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FILE COVERS 1907 - 10 Dec 2004 VOL 141 ISS 25  
FILE LAST UPDATED: 9 Dec 2004 (20041209/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> S L3/PREP

27 L3  
3233278 PREP/RL  
L4 25 L3/PREP  
(L3 (L) PREP/RL)

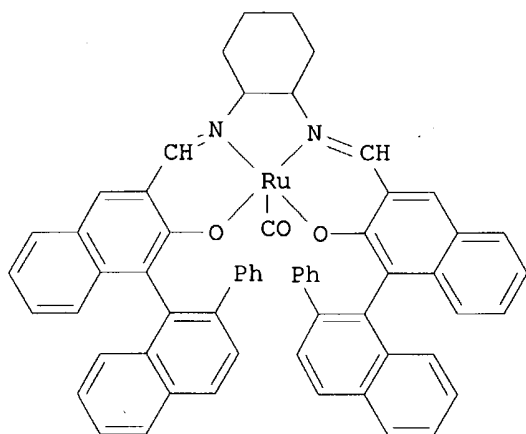
=> S L4 AND AZIDE

39169 AZIDE  
L5 7 L4 AND AZIDE

=> D 1-7 BIB ABS

L5 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2004:716150 CAPLUS  
DN 141:235107  
TI Method of producing optically active sulfimide compounds  
IN Katsuki, Tsutomu; Tamura, Yusuke; Uchida, Tatsuya  
PA Kyushu University, Japan  
SO Eur. Pat. Appl., 10 pp.  
CODEN: EPXXDW  
DT Patent  
LA English  
FAN.CNT 1

|      | PATENT NO.                                                                                                                        | KIND | DATE     | APPLICATION NO. | DATE     |
|------|-----------------------------------------------------------------------------------------------------------------------------------|------|----------|-----------------|----------|
| PI   | EP 1452523                                                                                                                        | A2   | 20040901 | EP 2004-250921  | 20040220 |
|      | EP 1452523                                                                                                                        | A3   | 20040929 |                 |          |
|      | R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK, HR |      |          |                 |          |
|      | JP 2004277398                                                                                                                     | A2   | 20041007 | JP 2003-310132  | 20030902 |
|      | US 2004230077                                                                                                                     | A1   | 20041118 | US 2004-777244  | 20040213 |
| PRAI | JP 2003-47744                                                                                                                     | A    | 20030225 |                 |          |
|      | JP 2003-310132                                                                                                                    | A    | 20030902 |                 |          |
| OS   | MARPAT 141:235107                                                                                                                 |      |          |                 |          |
| GI   |                                                                                                                                   |      |          |                 |          |



AB An optically active sulfimide compound is produced by using a specified Ru(salen)(CO) complex I as a catalyst by subjecting a specified alkyl aryl sulfide compound to an asym. sulfimidation with a specified **azide** compound having an easily eliminating group. For example, sulfimidation of PhSMe with N-butoxycarbonyl **azide** gave the corresponding optically active sulfimide in 18 % yield.

L5 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2003:262840 CAPLUS

DN 139:100713

TI Highly enantioselective (OC)Ru(salen)-catalyzed sulfimidation using N-alkoxycarbonyl **azide** as nitrene precursor

AU Tamura, Yuusuke; Uchida, Tatsuya; Katsuki, Tsutomu

CS Faculty of Science, Department of Chemistry, Kyushu University, Higashi-ku, Fukuoka, 812-8581, Japan

SO Tetrahedron Letters (2003), 44(16), 3301-3303

CODEN: TELEAY; ISSN: 0040-4039

PB Elsevier Science Ltd.

DT Journal

LA English

OS CASREACT 139:100713

AB Enantioselective imidation of alkyl aryl sulfides with N-alkoxycarbonyl **azide** as a nitrene precursor was effected by using (OC)Ru(salen) complex as catalyst. The steric and electronic nature of the N-alkoxycarbonyl group was found to strongly affect the enantioselectivity and the reaction rate, and high enantioselectivity (up to 99% ee) and good chemical yields were achieved by using 2,2,2-trichloro-1,1-dimethylethoxycarbonyl **azide** as the nitrene precursor at room temperature

RE.CNT 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2001:674578 CAPLUS

DN 136:102168

TI Ru(salen)-catalyzed asymmetric sulfimidation using arylsulfonyl **azide**

AU Murakami, M.; Uchida, T.; Katsuki, T.

CS Graduate School, Faculty of Science, Department of Chemistry, Kyushu University 33, CREST, JST (Japan Science and Technology), Hakozaki, Higashi-ku, Fukuoka, 812-8581, Japan

SO Tetrahedron Letters (2001), 42(40), 7071-7074

CODEN: TELEAY; ISSN: 0040-4039

PB Elsevier Science Ltd.

DT Journal  
LA English  
OS CASREACT 136:102168  
AB An (OC)Ru(II)(salen) complex was found to catalyze imidation of alkyl aryl sulfides in the presence of arylsulfonyl **azide** with high enantioselectivity as well as good chemical yield.

RE.CNT 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1999:447558 CAPLUS  
DN 131:214396  
TI The search for benchrotrenes and ferrocenes containing a chiral sulfoximido group. Preparation and structural properties  
AU Bolm, Carsten; Muniz, Kilian; Aguilar, Nuria; Kesselgruber, Martin; Raabe, Gerhard  
CS Institut Organische Chemie, RWTH, Aachen, D-52074, Germany  
SO Synthesis (1999), (7), 1251-1260  
CODEN: SYNTBF; ISSN: 0039-7881  
PB Georg Thieme Verlag  
DT Journal  
LA English  
OS CASREACT 131:214396  
AB Syntheses of chiral benchrotrene and ferrocene complexes bearing N-protected sulfonimidoyl moieties are reported. They were obtained by metal-catalyzed imination reactions starting from enantiopure sulfoxides. An x-ray structure anal. was carried out for one of the novel complexes confirming the stereospecificity of the imination process.

RE.CNT 62 THERE ARE 62 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1999:281299 CAPLUS  
DN 131:58629  
TI The preparation of N-tert-butyloxycarbonyl-(Boc-)protected sulfoximines and sulfimines by an iron(II)-mediated nitrene transfer from BocN3 to sulfoxides and sulfides  
AU Bach, Thorsten; Korber, Christina  
CS Fachbereich Chemie, Philipps-Univ., Marburg, D-35032, Germany  
SO European Journal of Organic Chemistry (1999), (5), 1033-1039  
CODEN: EJOCFK; ISSN: 1434-193X  
PB Wiley-VCH Verlag GmbH  
DT Journal  
LA English  
OS CASREACT 131:58629  
AB The imidation of sulfides and sulfoxides to the corresponding sulfimides and sulfoximides was carried out with Me3CO2CN3 (BocN3) in the presence of FeCl2. Sulfoxides reacted at room temperature in CH2Cl2 to give the corresponding sulfoximides in 40-95% yield. The imidation of the sterically congested Me3C(Me)S:O proceeded sluggishly (10% yield). The stereospecificity of the reaction was demonstrated with enantiomerically enriched (R)-(+)-PhMeS:O and (S)-(-)-PhCH2(Me)S:O which yielded the corresponding sulfoximides with retention of configuration. Mechanistically, an intermediate nitrene Fe(IV) complex is postulated as the reactive nitrene transfer reagent which is formed from FeCl2 and BocN3. The more nucleophilic sulfides reacted more readily in the imidation than sulfoxides. Their conversion to the corresponding sulfimides was conducted with BocN3 and 0.25 equiv FeCl2 in yields of 44-92%. In an alternative reaction mode, BocN3 was utilized at 0° in the presence of FeCl2 and Ac2CH2. The sulfimidation, which did not otherwise occur at this temperature, was accelerated by the ligand (36-90% yield).

RE.CNT 117 THERE ARE 117 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1998:435231 CAPLUS  
DN 129:148598  
TI Iron(II)-mediated nitrene transfer from t-butyloxycarbonyl azide  
(BocN3) to sulfoxides, sulfides, and ketene acetals  
AU Bach, Thorsten; Korber, Christina  
CS Fachbereich Chemie der Philipps-Universitat Marburg, Marburg, D-35032,  
Germany  
SO Tetrahedron Letters (1998), 39(28), 5015-5016  
CODEN: TELEAY; ISSN: 0040-4039  
PB Elsevier Science Ltd.  
DT Journal  
LA English  
OS CASREACT 129:148598  
AB The nitrene transfer from BocN3 to several nucleophiles is promoted by  
FeCl2 and yields the corresponding N-Boc protected sulfoximides,  
sulfimides, or  $\alpha$ -aminoalkanoates. Whereas the sulfoximide formation  
occurs spontaneously in CH2Cl2 as solvent the FeCl2-catalyzed nitrene  
transfer to sulfides and ketene acetals requires addition of a polar solvent.  
DMF was found to be best suited for this purpose.

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1972:72227 CAPLUS  
DN 76:72227  
TI Herbicidal and pesticidal sulfoximine derivatives  
IN Kirby, Peter; Whitaker, Kenneth E.  
PA Shell Internationale Research Maatschappij N. V.  
SO Ger. Offen., 35 pp.  
CODEN: GWXXBX  
DT Patent  
LA German

FAN.CNT 1

|      | PATENT NO.    | KIND | DATE     | APPLICATION NO. | DATE     |
|------|---------------|------|----------|-----------------|----------|
| PI   | DE 2129678    | A    | 19711230 | DE 1971-2129678 | 19710615 |
|      | GB 1307271    | A    | 19730214 | GB 1970-30895   | 19700625 |
|      | NL 7108165    | A    | 19711228 | NL 1971-8165    | 19710615 |
|      | FR 2096478    | A5   | 19720218 | FR 1971-21678   | 19710615 |
|      | FR 2096478    | B1   | 19740621 |                 |          |
|      | CH 554635     | A    | 19741015 | CH 1971-8737    | 19710615 |
| PRAI | GB 1970-30895 |      | 19700625 |                 |          |

AB Title compds. (.apprx.60) were prepared Thus, 11 g 2,4-Cl(O2N)-C6H3SOMe in  
200 ml CHCl2 and 20 ml H2SO4 was treated with 6.5 g NaN3 over 3 hr at  
48-55° to give 2,4-Cl(O2N)C6H3S(O)-(:NH)Me. p-ClC6H4S(O)(:NH)Me  
was treated with fuming H2SO4 and fuming HNO3 at 5-15°, and the  
mixture heated 6 hr at 30-5° to give 4,3-Cl(O2N)C6H3S(O)(:NH)Me.  
Biol. test data (herbicidal, acaricidal, insecticidal) were given.